

PATENT

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicant:

Allen, Keith D.

Examiner:

Wilson, Michael C.

Serial No.:

10/005,202

Group Art Unit:

1632

Filed:

12/04/2001

Docket No.:

R902/75658.023400

Title:

TRANSGENIC MICE CONTAINING KIR5.1 INWARDLY RECTIFYING

POTASSIUM CHANNEL GENE DISRUPTIONS

DECLARATION OF ROBERT DRISCOLL PURSUANT TO 37 C.F.R. § 1.132

Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

Sir:

I, Robert Driscoll, residing at 23 Chicory Lane, San Carlos, CA 94070, hereby declare:

- 1. I am presently employed as Vice President of Intellectual Property & Legal Affairs at Assignee, Deltagen, Inc., in San Carlos, CA. I have also previously served as the Company's Senior Director of Intellectual Property, in which position I managed and oversaw the Company's intellectual property portfolio, including the Company's patent filings. I possess a Ph.D in Chemistry, received from the California Institute of Technology. I also possess a J.D., received from Loyola Law School, Los Angeles. I am a registered patent attorney (Reg. No. 47,536).
- 2. I am familiar with the above-cited application. I am familiar with the Office Action mailed April 20, 2005. I am aware that the Examiner has rejected the claims, in part, for allegedly failing to meet the utility requirement. I am also aware that the Applicant has argued that a commercial sale of a mouse with a disrupted Kir5.1 inwardly rectifying potassium channel allele within the scope of the claimed subject matter ("Kir5.1 gene knockout mouse") should satisfy the utility requirement.

- 3. In support of the Applicant's aforementioned argument, I hereby state that I have reviewed Deltagen's internal sales records regarding the Kir5.1 gene knockout mouse. According to these records, the Kir5.1 gene knockout mouse has been delivered to at least one (1) large pharmaceutical company. The contractual terms by which the mice were transferred prohibit Deltagen from identifying the name of this company. However, the company is ranked among the top 10 pharmaceutical companies worldwide (based on sales).
- 4. It is my understanding, based on communications with our pharmaceutical company customers, that transgenic knockout mice obtained from Deltagen are used for studying gene function and for human therapeutic drug development.
- 5. I further declare that all statements made herein of my own knowledge are true; and further that these statements were made with knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the above-referenced application or any patent issuing thereon.

Robert Driscoll, Ph.D. Reg. No. 47,536

9 June 2

Date

PATENT

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Title: TRANSGENIC MICE CONTAINING KIR5.1 INWARDLY RECTIFYING

DECLARATION OF JOHN BURKE PURSUANT TO 37 C.F.R. § 1.132

POTASSIUM CHANNEL GENE DISRUPTIONS

Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

Sir:

I, John E. Burke, residing at 16357 E. Berry Avenue, Centennial CO 80015, hereby declare:

I am currently, and have been since 1998, the Attorney of Record for the 1. Applicant and Assignee, Deltagen, Inc. I am listed on the originally filed Power of Attorney for the present application. From December 1996 to December 1999, I was Of Counsel with the law firm of Pillsbury Madison & Sutro (currently Pillsbury Winthrop) where I represented Deltagen with respect to intellectual property matters, including patent matters relating to their transgenic mouse program. From December 1999 until December 2001, I served as Deltagen's Vice President of Intellectual Property, where I supervised Deltagen's internal patent department. All of the applications, including the present application, covering the 750 lines of mice in DeltaBase were drafted by Deltagen's patent department. From December 2001 until April 2003, I served as Deltagen's Senior Vice President and General Counsel. From April 2003 through April 2005, I was a partner with the Denver office of Merchant & Gould, where I continued to represent Deltagen with regard to intellectual matters, including patent matters. I am presently employed as a Shareholder with the Denver office of the law firm of Greenberg Traurig, where I am responsible for prosecution of Deltagens's patent portfolio relating to their transgenic mice program, including the present application.

- 2. I am familiar with the present application. I am familiar with the Office Action mailed April 20, 2005. I am aware that the Examiner has rejected the claims, in part, for allegedly failing to meet the utility and enablement requirements. I am aware that the Examiner argues on pages 7-8 of the Office Action that background and genetic factors have an effect on phenotypes such as the startle response.
- 3. I hereby declare that, as evidenced by the attached Exhibit, the subject matter of the present application, Kir5.1 inwardly rectifying potassium channel gene knockout mice (Kir5.1 gene knockout mice), were compared with control mice of identical background.
- 4. I hereby declare that the claimed Kir5.1 gene knockout mouse has been extensively analyzed using the tests set forth in the Examples. This data has been incorporated into Deltagen's commercial database product, DeltaBase. This database has been subscribed to by at least three of the world's largest pharmaceutical companies, Merck, Pfizer and GSK.
- 5. I hereby declare that I have accessed Deltagen's internal web-based DeltaBase database to review the data derived from analyses of the claimed mice. I hereby declare that the attached Exhibit contains four (4) pages, each representing a screen printout from DeltaBase. The first page is the Behavior Summary page summarizing changes relating to genotype associated with Gene 902, as prepared by Deltagen's pathology group. As noted at the top of the page, Gene 902 corresponds to the Kir5.1 gene. As noted, the homozygous mice were significantly different in their startle response relative to wild-type controls. The page further notes that for the behavioral tests, 9 homozygous mutant males were compared with 10 wild-type controls males. The page further describes how 129/OlaHsd x C57BL/6 F2N1 mice were produced. The table on the right side of the page provides the background of each of the homozygous (-/-) and wild-type control mice (+/+) used in the comparative tests. As is shown in the table, the transgenic mice (-/-) and control mice (+/+) are of identical F2N1 background (129/OlaHsd x C57BL/6).
- 6. The second and third pages represent the "left" and "right" sides of a webpage showing a portion of raw data derived from the comparative startle/PPI tests for Gene 902. The gene number, 902, is indicated in column 1. The ES cell line is indicated in column 2. As can be seen, each mouse was derived from the same ES cell line, 1006. Columns 3 and 4 indicate the

generation and background. As can be seen, each mouse is of identical background, F2N1. As can also be seen, each mouse tested is of approximately the same age and gender. Column 35 (page 3) shows the date on which the data was recorded. All of the data corresponding to the F2N1 mice was entered prior to the filing date of the present application.

- 7. The fourth page of the Exhibit shows the statistical values for the startle test for Gene 902. As shown, the data derived was statistically significant.
- 8. In summary, the attached Exhibits show that the transgenic mice were compared with control mice of identical background. The phenotypes were based on a comparison with age, gender and strain matched control mice.
- 9. I further declare that all statements made herein of my own knowledge are true; and further that these statements were made with knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the above-referenced application or any patent issuing thereon.

John E. Burke, Reg. No. 35,836

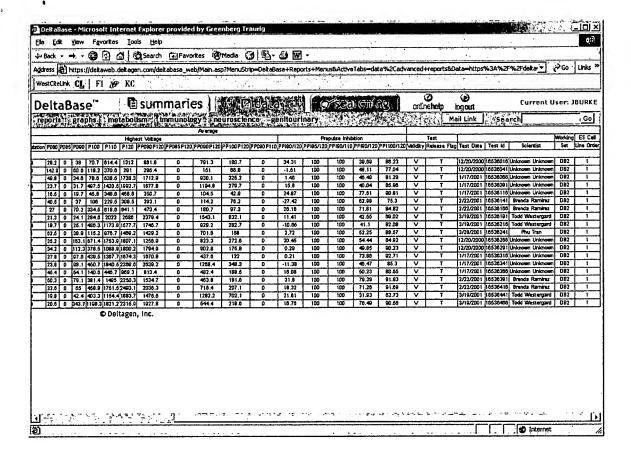
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gene orga	n system hot hits							Mail	Link	
Release: All F	Gene: 902 Name: Kcnj16 Family: Channel Subfamily: Potassiu	m <u>Al</u>	terna	tive Nam	es		1			1
Program: All F	Nucleotide Sequence Accession: AB016197 GI: 3953532 Ex	ternal	Links	Select	Exter	nal D	atab	ase i√	Ī	
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	Gene 902				Mic	ce				1
957 genes fou 🤌	_	#	Sex	Genotype	F	N	Age	Validity	Release	
9 Channel	Behavior	111500	Mala		_	 	Ť			1
10 Channel			_				├──			1
11 Channel	Changes related to genotype:		-		_		-			H
12 Channel			_			-			-	1
13 Channel ,	 Homozygous mutants displayed increased 		_		_	-	-		-	1
15 Protease	startle responses across all stimulus levels in					├-	_		 	l'
16 Protease Ir.	the Startle/PPI test.			-/-	2	1	_	v	T	1
17 GPCR		117320	Male	-/-	2	1	85	V	T	1
20 GPCR	Homozygous mutant and wild-type control mice	117341	Male	-/-	2	1	68	٧	T	1:
21 Carbohydri	were evaluated for phenotypic changes by testing	117341	Male	-/-	2	1	85	٧	Т	1
	on six behavioral tasks: Open field test, Tail	127734	Male	-/-	2	1	69	V	T	1
24 Carbohydri	suspension test, Rotarod test, Hot plate test,	127734	Male	-/-	2	1	84	٧	T]
25 Carbohydri	Startle/PPI, and Metrazol test.	127771	Male	-/-	2	1	68	٧	Т];
28 Carbohydri		Mail Link Mail	T].						
31 Carbohydri	Mouse ID numbers are as follows:	135164	## Alternative Names ## Sex Genotype Fen. Gen. Gen.]						
37 Growth Fa		135184	Male	-/-	2	1	70	V,	T	3
41 GPCR	9 homozygous mutant males (117341, 127734,	135184	Male	-/-	2	Mail Link Mail Mail Mail Mail Mail Mail Mail Mail				
<u>42</u> GPCR	127771, 135164, 111590, 117319, 117320, 111592,	111564	Male	+/+		1	_	٧	Т	
43 GPCR (▼	135184)	\vdash	-			1	-			
T. D	10 wild-type control males (138630, 127778,			i		1				
Phenotypic	117321, 117348, 117338, 135163, 111564, 111565,		-			<u> </u>	-			ĮÌ.
Release Summary	127772, 135180)	_				-	\vdash			-
Phenotypic					_	H-	-			1
<u>Data</u> Statistics	ES cells derived from the 129/OlaHsd mouse				-	H	_			ł
Target Research	substrain were used to generate chimeric mice.					\vdash	-			
Database	F1 mice were generated by breeding with				_	H	-			1
SUMMARY ANNOTATIONS	C57BL/6 females. The resultant F1N0				_	-	_			
H MOLECULAR BI	heterozygotes were backcrossed to C57BL/6 mice									
EXPRESSION AN EXPRESSION AN	to generate F1N1 heterozygotes. F2N1				-	\vdash	\vdash			
E CLINICAL FINDI	homozygous mutant mice were produced by	_		+/+	2	1	84	v		
⊞ BEHAVIOR	intercrossing F1N1 heterozygous males and	135163	Male	+/+	2	1	71	٧	Т	
# FERTILITY/DEV	formulas	135163	Male	+/+	2	1	80	٧	T	1
LE NEOROSCIENCE		135180	Male	+/+	2	1	70	٧	T	1
!	Behavior Findings:	135180	Male	+/+	2	1	79	٧	Т]:
;	When compared to age- and gender-matched	138630	Male	+/+	2	1	70	٧	Т	
į	wild-type control mice, homozygous mutant mice	138630	Male	+/+	2	1	84	٧	Τ	J
	were significantly different in their startle									
1	response on the Startle/PPI test. Mutants									1
	displayed higher startle responses than their									
	wildtype littermates. This may indicate a									
j	propensity for increased fear.									
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Del	taB	DeltaBase"				il summantes				(S)	C AV	G	36.5E	CF Sing	1000	© onlinehelp	3 brooks	_ 5	٥	urrent L	Current User: JBURKE	URKE
reports		graphs	meta	metabolism	imm	immunology	neur	neuroscience	uce	gen	genitourinary	Jary					MailsLink	link	Search		,	8
															Average							
ES	ES Cell							-				Highe	Highest Voltage						Prep	Prepulse Inhibition	, LO	
Gene	Line F#	N #Genotys	pe Gender	Age at Test	Mouse C	F #IN #Genotype GenderlAge at Test Mouse Chamber No Stimulation P080 P085 P09	timulation	2080 Po	85 P09	0 P100	P110	P120 P	P120 PP080P120 PP085P120 PP090P120 PP100P120 PP090P110 PP180/120 PP185/120 PP190/120 PP190/120 PP100/1	2085P120₽	P090P120	PP100P120P	P090P110	PP180/120F	P185/120P	P190/110P	P190/120P	P1100/12
				(days)																		
902	1006 2	+/+	Male	23	111564	1	42.4	28.2 0	0 38	70.7	614.4	1312	861.8	0	791.3	180.7	0	34.31	100	100	39.69	86.23
902	1006 2	+/+	Male	23	111565	2	70.8	142.9 0	0 60.6	118.2	370.6	291	295.4	0	151	8.99	0	-1.51	100	100	48.11	77.04
905	1006 2	+/+	Male	2.2	117321	8	48.7	49.8 0	34.6	9'8'8	538.5 1738.3	1738.3	1712.9	0	930.1	325.2	0	1.46	100	100	46.49	81.29
902	1006 2	1 +/+	Male	22	117338	1	24.6	23.7 0	0 31.7	7 497.5	1420.5 1992.7	1992.7	1677.8	0	1194.8	279.7	0	15.8	100	100	40.04	85.96
902	1006 2	1 +/+	Male	22	117348	3	25.6	16.5 0	0 19.7	7 45.8	348.6	466.8	350.7	0	104.5	42.9	0	24.87	100	100	77.61	90.81
902	1006 2	1 +/+	Male	22	127772	9	31.2	40.5 0		106	229.5	308.5	393.1	0	114.2	76.2	0	-27.42	100	100	62.98	75.3
902	1006 2	+/+	Male	82	127778	4	21.7	27 0	0 70.2	2 334.5	618.6	641.1	473.4	0	180.7	97.3	0	26.16	100	100	71.81	84.82
902	1006 2	1 +/+	Male	77	135163	1	23.2	21.3 0	0 24.	1 284.6	2023	2686	2379.4	0	1543.1	832.1	0	11.41	100	100	42.55	69.02
902	1006 2	1 +/+	Male	92	135180	3	16.4	19.7 0	0 25.1	1 486.3	3 1173.8 1577.7	1577.7	1745.7	0	929.2	282.7	0	-10.65	100	100	41.1	85.08
905	1006 2	+/+	Male	78	138630	5	51.1	62.5 0	38.8	9 115.2	976.7	1469.2	1429.2	0	701.6	168	0	2.72	100	100	52.25	88.57
902	1006 2	-/-	Male	73	111590	3	23.7	35.2 0	0 163.	Τ.	671.4 1753.9 1807.1	1807.1	1256.9	0	823.3	272.6	0	30.45	100	100	54.44	84.92
902	1006 2	·/	Male		111592	4	42.4	34.2 0	112.	.3 378.5	5 1088.91800.2	1800.2	1794.9	0	902.8	175.8	0	0.29	100	100	49.85	90.23
902	1006 2	1 -/-	Male	2.2	117319	9	37.5	27.8 0	0 97.6	6 439.5	439.5 1367.7 1674.3	1674.3	1670.8	0	437.6	122	0	0.21	100	100	73.86	92.71
902	1006 2	1 -/-	Male	22	117320	7	20.6	23.8 0	0 98.	1 450.7	7 1840.52369.6	2369.6	2639.2	0	1268.4	348.3	0	-11.38	100	100	46.47	85.3
902	1006 2	1 -/-	Male	2.2	117341	2	30.3	45.4 0	0 54.	1 140.6	446.7	969.3	813.4	0	482.4	188.5	0	16.08	100	100	50.23	80.55
902	1006 2	1 -/-	Male	82	127734	3	35.8	50.3	0 79.1	1 381.4	1495	2250.3	1534.7	0	463.8	181.6	0	31.8	100	100	79.39	91.93
905	1006 2	1 -/-	Male	77	127771	5	19.7	22.6 0) 25	458.9	1751.52493.	2493.1	2036.3	0	716.4	207.1	0	18.32	100	100	71.26	91.69
902	1006 2	1 -/-	Male	2.2	135164	2	16.8	19.8	0 42.4	#	403.3 1154.4 1883.7	1883.7	1476.6	0	1282.2	702.1	0	21.61	100	100	31.93	62.73
902	1006 2	1 -/-	Male	92	135184	4	30.8	20.6 0	343.	_	1198.31821.22315.9	315.9	1927.8	0	544.4	218.6	0	16.76	100	100	76.49	90.56
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	ES Cell						Study	·		St	tatistic			٠
Gene	Line	F#	N #	Age Bi	n Genotype	Gender	Procedure Name	Observable	Statistic	\	/alue	Mouse Count		
												(n)		
902	1006	2	1	49	-/-	Female	necropsy	liver weight	1-p value vs. WT co	ontrols	0.96	3		
902	1006	2	1	49	-/-	Female	necropsy weights	liver weight	1-p value vs. WT co	ontrols	0.96	3		
902	1006	2	1	90	-/-	Female	hematology	platelets	1-p value vs. WT co	ontrols	0.95	4		
902	1006	2	1	90	-/-	Male	serum chemistry	triglycerides	1-p value vs. WT co	ontrols	0.99	3		
902	1006	2	1	90	-/-	Male	serum chemistry	low density lipoprotein	1-p value vs. WT co	ontrols	0.99	3		
902	1006	2	1	180	-/-	Male	hematology	absolute basophils	1-p value vs. WT co	ontrols	0.97	4		
902	1006	2	1	180	-/-	Male	mouse metrics	body length	1-p value vs. WT co	ntrols	1	3		
902	1006		1		-/-	Male	startle	P090, average	1-p value vs. WT co	ontrols	0.98	9		
902	1006		1		-/-	Male	startle	P090, standard deviation	1-p value vs. WT co	ontrols	0.98	9		
902	1006		1		-/-	Male	startle	P100, average	1-p value vs. WT co	ontrols	0.98	9		
902	1006		1		-/-	Male	startle	P100, standard deviation	1-p value vs. WT co	ntrols	0.99	9		
902	1006		1		-/-	Male	startle	P110, average	1-p value vs. WT co	ontrols	0.97	9		
902	1006		1		-/-	Male	startle	P110, standard deviation	1-p value vs. WT co	ontrols	0.97	9		
902	1006		1		-/-	Male	startle	P120, average	1-p value vs. WT co	ntrols	0.97	9		
902	1006		1		-/-	Male	startle	P120, standard deviation	1-p value vs. WT co	ntrols	0.96	9		

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